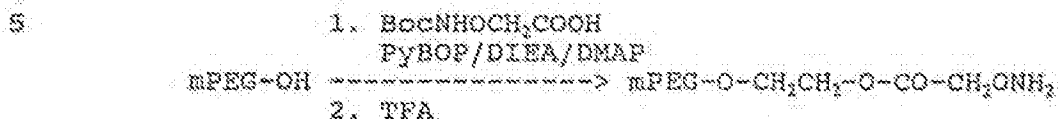


treated with cold ether. A white product was obtained. Yield 0.5g. IR: (C=O): 1676. Analysis. Calcd. for N, 0.55. Found: N, 0.50.

Synthesis of $\text{CH}_3\text{O}-(\text{CH}_2\text{CH}_2\text{O})_n-\text{CH}_2\text{CH}_2-\text{O}-\text{CO}-\text{CH}_2-\text{ONH}_2$



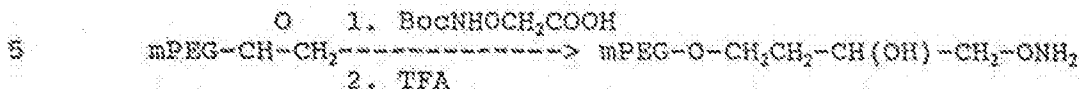
10 mPEG-5000-alcohol (2.0g, 0.4 mmol) and Boc-aminooxyacetic acid (0.2g, 1.05 mmol) were dissolved in 20 ml dichloromethane. Benzotriazol-1-yl-oxytripyrrolidinophosphonium hexafluorophosphate (1.1g, 2mmol) was added followed by diisopropylethylamine (0.7ml, 3.09 mmol). The reaction was stirred for about 2 hours at room temperature after which time dimethylaminopyridine (0.244g, 2 mmol) was added. After about four days cold ether was added, and the resulting precipitate was collected by filtration, washed, and dried. The precipitate was taken up in water and was treated with decolorizing carbon. After about 24 hr. the decolorizing carbon was filtered, and the clear filtrate was concentrated. The residue was dissolved in dichloromethane, dried with sodium sulfate, filtered, and the filtrate was treated with cold ether. Analysis. Calcd. for N, 0.28. Found: N, 0.14.

25 Half of the collected precipitate was placed in a dichloromethane/trifluoroacetic acid (2:1) mixture. After 30 minutes the solution was concentrated. The compound was further purified by gel filtration on a

30

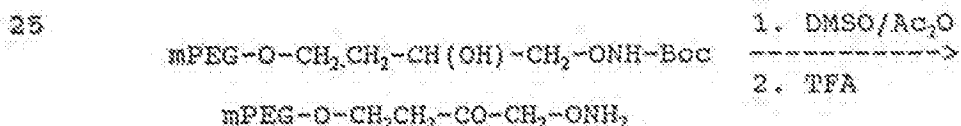
LH-20 column eluting with methanol/methylene chloride (5:1). Yield 0.3g. IR: (C=O): 1734.

Synthesis of CH₃O-(CH₂CH₂O)_n-CH₂CH₂-CH(OH)-CH₂-ONH₂



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mPEG-5000-epoxide (1.0g, 0.2 mmol) was dissolved in 10ml 0.1M NaOH. t-Butyl N-hydroxycarbamate (0.53g, 4 mmol) was added. After running the reaction overnight the reaction mixture was extracted with dichloromethane. Sodium sulfate was added and was filtered. Cold ether was added the dichloromethane solution, and the resulting precipitate was collected by filtration, washed, and dried. The compound was further purified by gel filtration on a LH-20 column eluting with methanol/methylene chloride (5:1). The protected mPEG-derivative (0.25g) was placed in a dichloromethane/trifluoroacetic acid (1:1) mixture. After 30 minutes the solution was concentrated and taken up in dichloromethane. The compound was isolated by precipitation from ether. Yield 0.2g. IR: (O-H): 3447. Analysis. Calcd. for N, 0.28. Found: N, 0.21.

Synthesis of CH₃O-(CH₂CH₂O)_n-CH₂CH₂-CO-CH₂-ONH₂



mPEG-O-CH₂CH₂-CH(OH)-CH₂-ONH-Boc (0.3g, 0.6 mmol) was placed in 3 ml dry dimethyl sulfoxide followed by the addition of 2 ml dry acetic anhydride. The reaction went for about 24 hr. at room temperature after which time cold ether was added. The resulting precipitate was collected by filtration, washed and dried. The product was placed in a dichloromethane/trifluoroacetic acid (1:1) mixture. After 30 minutes the solution was concentrated. The compound was isolated by precipitation from cold ether. Yield 0.18g. IR: (C=O): 1698. Analysis. Calcd. for N, 0.28. Found: N, 0.20.

Synthesis of mPEG-Ornithine Semicarbazide

mPEG-5000-amine (5.0 g, 1. mmol) was dissolved in 5 ml dry methylene chloride and 10 ml dry dimethylformamide was added. Fmoc-Orn(Boc)-OPfp (3.1 g, 5 mmole) was added. After 1 hr. at room temperature the solution was concentrated. Water was added to the residue, and the solution was filtered, centrifuged, and filtered to remove the dispersed solid in the aqueous solution. The filtered aqueous solution was concentrated, and the residue was taken up in methylene chloride and was dried with sodium sulfate. The solution was filtered. The filtrate was treated with cold ether. The resulting precipitate was collected by filtration, washed, and dried. Yield 3.3 g. IR: (C=O):1713, 1680. The Fmoc group was removed by treating the compound with 25% piperidine (in methylene chloride) for 30 min. Cold ether was added to the solution to precipitate the mPEG-derivative. The precipitate was collected, washed, and dried. The free alpha amino group was acetylated by dissolving 1.4 g of the mPEG-derivative

with 3 ml methylene chloride and adding 1 ml acetic
anhydride. After about 1.7 hr. at room temperature
the solution was concentrated. The resulting solid
was treated with trifluoroacetic acid in methylene
chloride (3:5) for 1 hr. at room temperature. The
solvent was removed and resulting oil was taken up in
methylene chloride and cold ether was added. The
precipitate formed was collected, washed, and dried.
Yield 1.1 g. IR: (C=O): 1685. The mPEG-derivative
was dissolved in 3 ml dry methylene chloride and
triethylamine (0.54 ml, 3.84 mmole) was added
followed by 1 ml phosgene in toluene (1.92 mmole) and
an additional 2 ml dry methylene chloride. The
reaction went overnight at room temperature after
which time the solvent was removed. The residue was
dissolved in 3 ml dry methylene chloride and 0.2 ml
hydrazine (5.75 mmole) was added. Dry methanol was
added until the solution became clear (3.4 ml).
After 4 hr. at room temperature the solution was
clarified by centrifugation and was concentrated.
The compound was purified by gel filtration on a LH-
20 column eluting with methanol/methylene chloride
(5:1). Yield 0.7 g. IR: (C=O): 1675.

The invention having been described, the
following examples are offered by way of
illustration, not by way of limitation, of the
subject invention.

EXAMPLES

Modification of EPO (Hydrazide Method)

In a typical experiment, EPO (0.5-1.0 mg)
(obtained from Ortho Biotech) was placed in 100 mM
sodium acetate, pH 5.6, total volume 0.786 ml.

Enough 10 mg/ml solution of sodium periodate was added to give a final concentration of sodium periodate at 10 mMol. The oxidation went for 30 min at 0°C in the dark after which time 0.33 ml 80 mMol Na₂SO₃ was added. After 5 min the solution was concentrated and washed three times with 100 mMol sodium acetate, pH 4.2 in a microconcentrator. After the final concentration the oxidized EPO solution was brought up to 1.0 ml with 100 mM sodium acetate. mPEG5000-hydrazide (50 mg) was added to the oxidized EPO. The mixture was stirred over night at room temperature. The mPEG5000-EPO was purified by gel filtration using a Sephacryl S-200-HR column (1 mm x 45 mm) eluting with a phosphate buffer containing 0.05% sodium azide. The amount of mPEG modifying EPO was determined by HPLC gel filtration using either a Zorbax® GF-250 or GF-450 column using a 0.1 M phosphate buffer, pH 7. From 6-12 molecules mPEGs were found to be attached to each molecule of EPO.

20 Modification of EPO (Semicarbazide Method)

The same procedure as above for the hydrazide method was performed, except the reaction time for oxidation was decreased to 5 minutes and a decreased amount of mPEG-semicarbazide (10 mg) was used compared to the amount of mPEG-hydrazide (50 mg). Even with the decreased oxidation time and less mPEG added, more (about 18) mPEG molecules were attached to EPO. If longer oxidation times (15 min) and more mPEG-semicarbazide is added, around 30 mPEG molecules can be attached to EPO depending on the molecular weight of mPEG used. Thus mPEG-semicarbazide appears to be more reactive than mPEG-hydrazide and attaches

many more mPEG molecules to EPO than mPEG-hydrazide is able to under similar reaction conditions.

A comparison of the effect of modifying EPO with mPEG on either the carbohydrate groups or on the amino acid side chains is shown in Fig. 1.

5 Analytical HPLC gel filtration conditions are the same as described above. The chromatogram of unmodified EPO is presented in Fig. 1a. A single peak with a retention time of 10.5 min is found.

10 When EPO is modified with mPEG5000 on its carbohydrate groups (Fig. 1b), a single large peak with a retention time of 9.4 minutes is seen for the unpurified reaction product. When EPO is reacted with a succinimide ester of mPEG5000 which reacts with the side chain of lysine, a heterogeneous mixture of reaction products is obtained (peaks from 7.5 - 10.2 minutes). A similar heterogeneous pattern for mPEG modification using succinimide coupling to CSF-1, interleukin-2, and β -interferon has been found (U.S. Pat No. 4,847,325 and 4,9117,888). There are also more low molecular weight impurities present with succinimide coupling (Fig. 1c). Active ester coupling using succinimide derivatives of mPEG has been the preferred method for attaching mPEG to

15 proteins [Nucci, M.L., Shorr, E., and Abuchowski, A. (1991) Adv. Drug Delivery Rev., 5, 133-151]. EPO was also derivatized with mPEG8500 using the above-described semicarbazide method, see Figure 4 for results of biological experiments. The above-

20 described semicarbazide method was also used to obtain EPO modified with mPEG12000 (see Figure 3) and EPO modified with mPEG2000 (see Figure 10).

EPO also was modified with thiosemicarbazide, hydrazide carboxylate, and carbonic acid dihydrazide

derivatives of mPEG. These derivatives of mPEG performed like the semicarbazide derivatives of mPEG in that high levels of coupling mPEG to EPO could be obtained using these derivatives when compared to the hydrazide derivatives of mPEG. Other conditions for the oxidation of EPO can be used such as increased temperature, increased concentration of sodium periodate, and increased or decreased reaction times as long as these oxidation conditions do not impair the biological activity of EPO.

Large-Scale Modification of EPO (Semicarbazide or Carboxylate Hydrazide Method)

EPO (12.0mg) (obtained from Ortho Biotech) was placed in 100 mM sodium acetate, pH 5.5, total volume 1.8 ml. Sodium periodate (0.215 ml) at a concentration of 40 mg/ml was added. The oxidation went for 20 minutes at 0°C in the dark after which time 0.02 ml of ethylene glycol was added, and the admixture stirred for 10 minutes at 0°C. The oxidized-EPO was purified by gel filtration using a Sephadex® G-25 column (2.5 cm X 9 cm) and eluted with 100mM sodium acetate buffer, pH 4.3. Eluted oxidized EPO (10-11ml) was pooled. mPEG5000 semicarbazide (100 mg) was added to the purified oxidized EPO. The mixture was stirred overnight at room temperature. The mPEG-5000 EPO was purified by gel filtration using a Sephacryl® S-200-HR column eluting with buffer consisting of 0.2M NaCl, 0.02 M sodium citrate, 0.025% sodium azide, pH 7.0.

The above modification was repeated using 200 mg of mPEG5000 semicarbazide. Reactivity was about 22 mPEG molecules per molecule of EPO.

The above modification was repeated employing half the amount of the reactants as specified

hereinabove, using 200 mg of carboxylate hydrazide. Reactivity was about 30 mPEG molecules per molecule of EPO.

Modification of EPO (Oxime Method)

5 A. mPEG-CH₂CH₂-NH-CO-CH₂-ONH₂

The same procedure as above for the large-scale semicarbazide method was performed. However, instead of addition of mPEG5000 semicarbazide, mPEG5000-CH₂-CH₂-NH-CO-CH₂-ONH₂ (50 mg) was admixed with 2.15 ml of oxidized EPO at room temperature overnight. The mPEG5000-EPO was purified by gel filtration using a Sephacryl® S-200-HR column eluting with buffer consisting of 0.2M NaCl, 0.02 M sodium citrate, 0.025% sodium azide, pH 7. About 31 molecules of mPEG5000 were found to be attached to each molecule of EPO as determined by HPLC gel filtration using a Phenomenex Biosep-Sec-S4000 column (30 cm X .917 cm). A minor fraction consisting of 25 molecules of mPEG-EPO was also isolated, and was used for biological testing. See Figures 16, 18 and 19.

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B. mPEG-O-CH₂CH₂-NH-CO-ONH₂

The above modification of EPO was repeated using mPEG-O-CH₂CH₂-NH-CO-ONH₂. About 18-19 molecules of mPEG5000 were found to be attached to each molecule of PEG as determined using the methods as in modification A above. Biological data is shown in Figures 16, 18 and 19.

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C. mPEG-O-CH₂CH₂-ONH₂

The above modification of EPO was repeated using mPEG-O-CH₂CH₂-ONH₂. About 17 molecules of mPEG5000 were found to be attached to each molecule of PEG as

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determined using the methods as in modification A above. Two minor fractions of 23mPEG, 12mPEG were also isolated(Figure 17, 18, 19).

D. mPEG-O-CH₂CH₂-CO-ONH₂

5 The above modification of EPO is repeated using the PEG-oxime derivative mPEG-O-CH₂CH₂-CO-ONH₂. About 3 molecules of mPEG5000 were found to be attached to each molecule of EPO, as determined using the methods as for modification A above.

10 E. mPEG-O-CH₂CH₂-CH(OH)-CH₂-ONH₂

The above modification of EPO is repeated using the PEG-oxime derivative mPEG-O-CH₂CH₂-CH(OH)-CH₂-ONH₂. About 31 molecules of mPEG5000 were found to be attached to each molecule of EPO, as determined using the methods as for modification A above.

15 F. mPEG-O-CH₂CH₂-NH-CS-ONH₂

The above modification of EPO is repeated using the PEG-oxime derivative mPEG-O-CH₂CH₂-NH-CS-ONH₂. About 4 molecules of mPEG5000 were found to be attached to each molecule of EPO, as determined using the methods as for modification A above.

Biological Activity of mPEG-EPO

25 The mPEG-EPO derivatives were assayed for biological activity in vivo by measuring the increase in erythrocytes generated after injection of the modified protein (Egrie, J.C., Strickland, T.W., Lane, J., Aoki, K., Cohen, A.M., Smalling, R., Trail, G., Lin, F.K., Browne, J.K., and Hines, D.K. (1986) Immunobiol. 172: 213-224). Briefly, mice (female CD-1, eight weeks old) were injected either

intraperitoneally or subcutaneously with 0.4 μ g protein once a day for two consecutive days. Blood was withdrawn on predetermined days for hematocrit readings.

5 The in vivo biological activities (hematocrit levels) of the mPEG-EPOs linked via hydrazide derivatives of mPEG is presented in Figures 2, 4 and 5 and Tables I and II. To summarize the findings, Figures 2 and 5 show that the optimal number of mPEG
10 coupling is not obvious and has to be determined by synthesis and biological testing in order to give the best mPEG-EPO. Figure 4 compares mPEG coupling using hydrazide and semicarbazide linkers. Higher and longer hematocrit levels for mPEG-EPO could be
15 obtained using the semicarbazide linker. The observed results are due, in part, to the higher level of incorporation of mPEG which could be obtained using the semicarbazide linker. Table I shows the biological activity of the mPEG-EPOs as a
20 function of the number of mPEGs incorporated and the molecular weight of mPEG used.

 Table I summarizes the biological activity of different hydrazide mPEG-EPOs comparing molecular weight of mPEG used and the amount of mPEG coupled.

25 Table II shows a comparison of the different mPEG hydrazide linkers used. Table II summarizes the biological activities of EPO modified with different mPEG5000-hydrazide derivatives where optimal amounts of mPEG were incorporated. Not all the carbohydrate mPEG-derivatives give the same biological activity
30 when coupled to EPO due to the inability to sufficiently couple an optimal amount of mPEG or other factors. Biological activities are the best values obtained for each linker.

TABLE I

Table I: Biological Activity of Different mPEG-EPOs

<u>#mPEGs Coupled *</u>	<u>Molecular Weight mPEG</u>	<u>Max. Hematocrit (%)**</u>	<u>Duration of Activity Relative to EPO (Days)***</u>
16	2000	60	14
8 #	5000	63	8
12 #	5000	61	7
18	5000	61	18
22	5000	61	20
24	5000	58	13
28	5000	54	14
17	8000	58	16
12 #	8500	54	7
20	8500	58	14
34	8500	47	7
8 #	12000	54	7
14	12000	53	11
28	12000	51	7

* As Determined by Size Exclusion Chromatography

** Biological Assay Described in Experimental Section

*** Days from EPO Maximum Hematocrit Level (48 on Day 4) Required to Reach EPO's Maximum Hematocrit Level After Attaining its Own Maximum Hematocrit Level

Hydrazide Linker Used. All Others Used Semicarbazide Linker.

Table II: Biological Activity of mPEG5000-EPOs Comparing Different Carbohydrate Modifying mPEG-Linkers

mPEG-Linker		Max. Hematocrit (%) *	Duration of Activity Relative to EPO (Days)**
CH ₃ -(OCH ₂ CH ₂) ₆ -D-CH ₂ -CO-NHNH ₂	(Hydrazide)	54	8
CH ₃ -(OCH ₂ CH ₂) ₆ -NH-CO-NHNH ₂	(Semicarbazide)	81	20
CH ₃ -(OCH ₂ CH ₂) ₆ -NH-CS-NR ₂	(Thiosemicarbazide)	59	14
CH ₃ -(OCH ₂ CH ₂) ₆ -NH-CO-NHNH-CO-NHNH ₂	(Carbonyl Acid Dihydrazide)	57	13
CH ₃ -(OCH ₂ CH ₂) ₆ -D-CO-NHNH ₂	(Hydrazide Carboxylete, 22 mPEGs)	60	
CH ₃ -(OCH ₂ CH ₂) ₆ -NH-CO-C ₆ H ₄ -NHNH ₂	(Aryhydrazide)	52	7

* Biological Assay Described in Experimental Section

** Days from EPO Maximum Hematocrit Level (40% on Day 4) Required to Reach EPO's Maximum Hematocrit Level After Attaining Its Own Maximum Hematocrit Level

The hematocrit levels for EPO and mPEG5000-EPOs in mice are presented in Fig. 2. The 12PEG-EPO was made by coupling mPEG5000-hydrazide and reflects the maximum incorporation which could be achieved by this mPEG derivative under the experimental conditions given above. The 18PEG and 28PEG EPOs were made by coupling with mPEG5000-semicarbazide. The semicarbazide derivatives of mPEG result in much better biological activity than the hydrazide derivatives of mPEG due to the larger amounts of mPEG which can be incorporated using this mPEG derivative. All three mPEG5000-EPOs show increased maximum and prolonged activity when compared to native EPO. Thus modification of a protein's carbohydrate groups with PEG can yield a much more potent therapeutic protein.

For additional data on the effects of various mPEG hydrazide-modified EPOs employed in the experiments on hematocrit levels, see Figures 8-15. The mPEG modified EPO employed in the experiments depicted in Figures 8-15 were prepared using the appropriate water-soluble polymer reagent essentially as described for the other mPEG modified EPO molecules used the experiments depicted in Figures 2-5.

The in vivo biological activities (hematocrit levels) of the mPEG-EPOs linked via oxime-forming derivatives of mPEG is presented in Figures 16 and 17. To summarize, Figure 16 compares mPEG coupling using mPEG-O-CH₂CH₂-NH-CO-ONH₂ ("A") and mPEG-O-CH₂CH₂-NH-CO-CH₂-ONH₂ ("C") linkers. Higher hematocrit levels of mPEG-EPO could be obtained using the "A" linker (corresponding to Formula XXX herein) having 18 mPEG molecules per molecule of EPO as compared to the "C" linker (corresponding to Formula

XXXIII herein) having 31 mPEG molecules per molecule of EPO. Also notable was the higher hematocrit activity of the "C" linker (Formula XXXIII herein) having 25 mPEG molecules per molecule of EPO as compared to the same linker having 31 molecules of mPEG per molecule of EPO.

Figure 17 compares mPEG coupling using mPEG-O-CH₂CH₂-ONH₂ (formula XXIII) ("B") linker at 22, 17, and 12 mPEG molecules per molecule of EPO. Highest hematocrit levels are obtained at the lowest degree of pegylation, and hematocrit was decreased inversely proportional to degree of pegylation. In all mPEG linkers using mPEG-O-CH₂CH₂-ONH₂ oxime derivative however, hematocrits were higher and of increased duration as compared to native EPO.

Especially noteworthy is the biological activity of the 12mPEG-EPO of formula XXIII. Hydrazide derivitized 12 mPEG-EPO produces neither the degree nor duration of hematocrit elevation as that of the oxylamine derivitized EPO.

Antibody binding to mPEG-EPO

The antigenicity of the mPEG-EPOs was determined by using the Clinigen™ erythropoietin (EPO) EIA test kit. Briefly, the assay consists of a micro titre plate coated with a monoclonal antibody to EPO. EPO or mPEG-EPO is allowed to interact with the coated plate. After washing the plate a labeled polyclonal antibody to EPO is incubated on the plate. After substrate development the plate is read.

The results of the ELISA assay for hydrazide derivitized EPO, are presented in Figure 3. The mPEG-EPOs are presented as the approximate number of mPEGs attached for a given molecular weight of mPEG.

For example, 12PEG-5k means about 12 mPEG molecules of a molecular weight of about 5000 were coupled to each molecule of EPO. The data indicates that as the number of mPEGs coupled to EPO are increased, the antigenicity of the protein is decreased. Similarly as the molecular weight of mPEG is increased, the antigenicity, i.e. the binding of the antibody, of the modified EPO also is decreased. Reacting oxidized EPO with a hydrazide derivative of mPEG did not reach the high coupling levels seen with semicarbazide, thiosemicarbazide, and carbonic acid dihydrazide PEG derivatives and thus could not give the large decreases in immunogenicity as seen with these other mPEG derivatives. Decreasing the antigenicity of a protein correlates to a decrease in the immunogenicity of a protein as well. Thus mPEG-EPO coupled to the carbohydrate groups of EPO may reduce any potential immunogenicity related to the protein with those derivatives of mPEG able to be coupled at high levels being the most effective.

For additional ELISA data with mPEG hydrazide derivatives see Figure 6.

The results of the ELISA assay for oxime derivatized EPO are presented in Figure 18. The data indicates that as the number of mPEGs coupled to EPO are increased, the antigenicity of the protein is decreased. Reacting oxidized EPO with a linker that resulted in comparatively low coupling levels (18 PEG-A, 17 PEG-B, 12 PEG-B) did not give the large decreases in immunogenicity as seen with the comparatively high coupling level formulations (32 PEG-B, 25 PEG-C, and 31 PEG-C. Note that these differences in a linker's ability to decrease immunogenicity appear to be determined largely based

on the coupling level (e.g. compare 12 PEG-B and 22PEG-B). Thus, mPEG coupled to carbohydrate groups of EPO through oxime linkages may reduce potential immunogenicity related to the protein, with those derivatives of mPEG able to be coupled at high levels being the most effective.

Modification of Horseradish Peroxidase: Comparison of Hydrazide Versus Semicarbazide Coupling

Horseradish peroxidase (HRP) is a glycoprotein enzyme (oxido-reductase). HRP was modified with either mPEG5000-hydrazide or mPEG5000-semicarbazide in order to see whether another glycoprotein besides EPO could show the difference in modification between the two different carbohydrate modification reagents. In a typical experiment, horseradish peroxidase (2 mg) was placed in 100 mM sodium acetate, pH 5.6, total volume 0.8 ml. Enough 10 mg/ml solution of sodium periodate was added to give a final concentration of sodium periodate at 10 mMol. The oxidation went for 15 min at 0°C in the dark, after which time 0.33 ml 80 mMol Na2SO3 was added. After 5 minutes, the solution was concentrated and washed three times with 100 mMol sodium acetate, pH 4.2, in a micro concentrator. The oxidized horseradish peroxidase solution was then split in half, with one half receiving mg PEG5000-hydrazide and the other half receiving 30 mg mPEG5000-semicarbazide. The two oxidized horseradish peroxidase solutions were then stirred overnight at room temperature. The extent of PEG modification was determined by HPLC gel filtration using a Zorbax™ CF-250 column using a 0.1 M phosphate buffer, pH 7. The mPEG5000-hydrazide modified horseradish peroxidase had approximately 7

PEG molecules/HRP; the mPEG5000-semicarbazide modified horseradish peroxidase had approximately 19 PEG molecules/HRP. Thus modification of horseradish peroxidase with PEG being attached to its carbohydrate groups is more effective using a semicarbazide derivative of PEG than a hydrazide under the same experimental conditions.

Half-Life Determinations

Half-Life experiments were done in male Sprague-Dawley rats weighing about 0.3 kg. Three rats were used for each compound. The experimental details are as follows. Each rat was injected IV (intravenously) with 1 μ g EPO or mPEG-EPO. For the hydrazide derivatized mPEG-EPO, the mPEG-EPO used was mPEG500 semicarbazide-18, prepared essentially as described in the section above on EPO modification with semicarbazides. Blood was withdrawn from each rat at 2, 5, 15, 45, 90 minutes and 3, 6, 24, 48, 54 hour time points. The blood was collected in heparinized tubes, and the plasma was isolated. The isolated plasma was tested for EPO biological activity in an EPO dependent cell proliferation assay. The in vitro assay employed used an FDC-P1/ER cell line. This murine cell line incorporates the EPO receptor and is dependent on EPO for growth. The assay was performed as follows. The cells were grown (10⁶/ml) in the absence of EPO for 24 hr after which time either EPO or mPEG-EPO at different concentrations is added to the cells. The cells were incubated for 42 hr, and then tritiated thymidine was added to the cells. After 6 hr the cells were harvested and counted. Cell growth was determined by the increased up-take of thymidine. Results are given in figure 7.

The EPO dependant cell proliferation assay was performed as described above using oxime-derivatized mPEG-EPO. Results are given in Figure 19.

In Vivo Assay: Anemic Mouse Model

9 In this assay mice are rendered anemic by injections for five consecutive days with TNF-alpha. To overcome the anemia the mice were injected SC (subcutaneously) with either EPO or mPEG-EPO (at 0.03 µg/dose) over the same five days or on just two of
10 the five days that the mice receive TNF-alpha. Results are given in figure 15.

Equivalents

All publications and patents mentioned in the above specification are herein incorporated by
15 reference. The foregoing written specification is considered to be sufficient to enable one skilled in the art to practice the invention. Indeed, various modifications of the above-described modes for carrying out the invention which are obvious to those
20 skilled in the field of molecular biology or related fields are intended to be within the scope of the following claims.

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:-

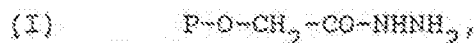
1. A compound having the formula:



5 wherein X is O or S; Q is selected from the group consisting of -NHNH₂, and -C₆H₄-NHNH₂; and Y is selected from the group consisting of -O-, -OCH₂-, -NH-, -NHNH-, -O-CO-CH₂CH₂- and -NHCO-N-NHNH-; and P is a water-soluble polymer.

10 2. A compound according to Claim 1, said compound belonging to the group consisting of,

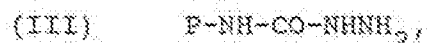
a compound having the formula:



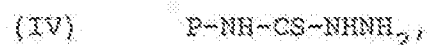
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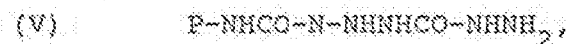
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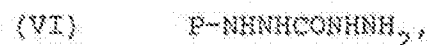
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20 a compound having the formula:



a compound having the formula:



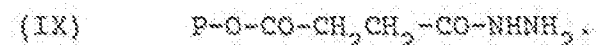
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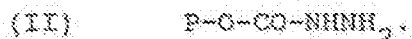
a compound having the formula:



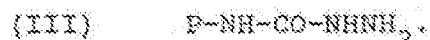
a compound having the formula:



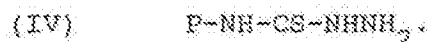
3. A compound according to Claim 2, said compound having the formula:



4. A compound according to Claim 2, said compound having the formula:



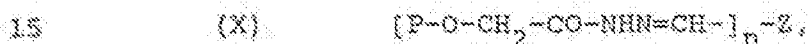
5. A compound according to Claim 2, said compound having the formula:



10 6. A compound according to Claim 2, said compound having the formula:



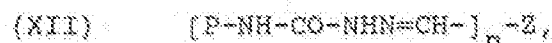
7. A compound belonging to the group consisting of, a compound having the formula:



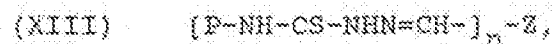
a compound having the formula:



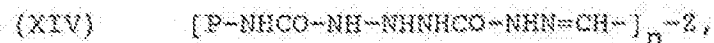
a compound having the formula:



20 a compound having the formula:



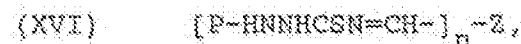
a compound having the formula:



a compound having the formula:



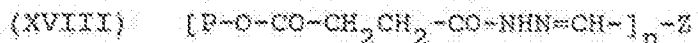
a compound having the formula:



a compound having the formula:

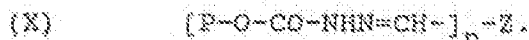


a compound having the formula:



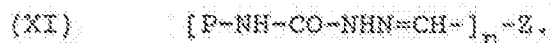
wherein Z is a polypeptide, n is 1 to x, x being the number of oxidation activatable groups on Z, and P is a water-soluble polymer.

8. A compound according to Claim 7, having the formula



9. The compound according to Claim 8 wherein n is 22-32.

10. A compound according to Claim 7, having the formula

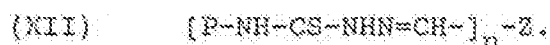


11. The compound according to Claim 10, wherein n is 10-36.

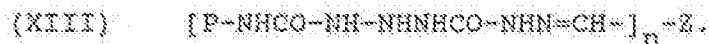
12. The compound according to Claim 10, wherein n is 20-32.

13. The compound according to Claim 10 wherein n is 17-25.

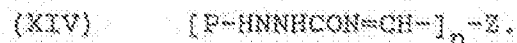
14. A compound according to Claim 7, having the formula



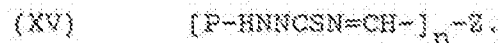
15. A compound according to Claim 7, having the formula



16. A compound according to Claim 7, having the formula



17. A compound according to Claim 7, having the formula



18. A compound having the formula:

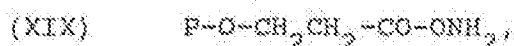


wherein X is C=O, S, CH₂ or CHOH; Q is selected from

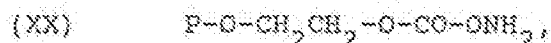
the group consisting of $-\text{ONH}_2-$ and $-\text{CH}_2-\text{ONH}_2-$, and Y is selected from the group consisting of $-\text{O}-\text{CH}_2\text{CH}_2-$, $-\text{O}-\text{CH}_2\text{CH}_2-\text{O}-$, $-\text{O}-\text{CH}_2\text{CH}_2-\text{N}-$, $\text{O}-\text{CH}_2\text{CH}_2-\text{S}$, and $-\text{O}-\text{CH}_2\text{CH}_2\text{CH}-$; and P is a water soluble polymer.

19. A compound according to Claim 18, said compound belonging to the group consisting of,

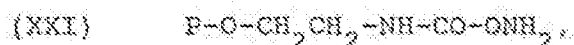
a compound having the formula:



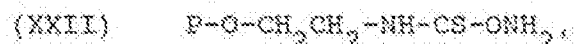
a compound having the formula:



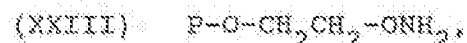
a compound having the formula:



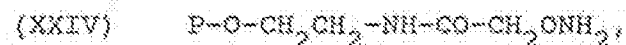
a compound having the formula:



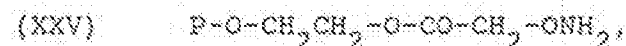
a compound having the formula:



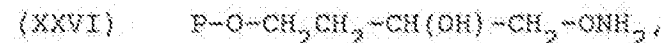
a compound having the formula:



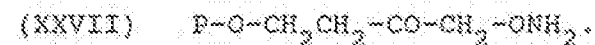
a compound having the formula:



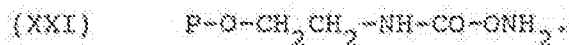
a compound having the formula:



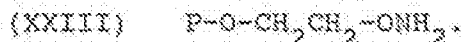
a compound having the formula:



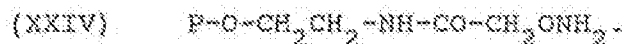
20. A compound according to Claim 19, said compound having the formula:



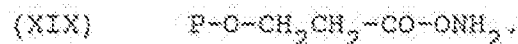
21. A compound according to Claim 19, said compound having the formula:



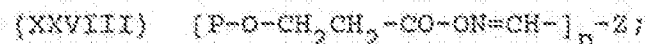
22. A compound according to Claim 19, said compound having the formula:



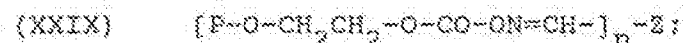
23. A compound according to Claim 19, said compound having the formula:



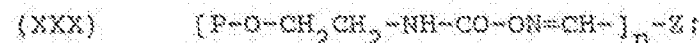
24. A compound belonging to the group consisting of, a compound having the formula:



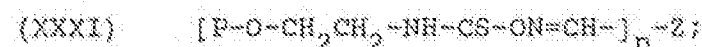
a compound having the formula:



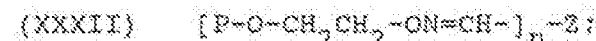
a compound having the formula:



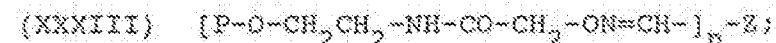
a compound having the formula:



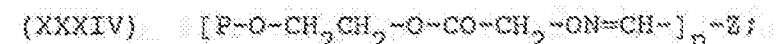
a compound having the formula:



a compound having the formula:



a compound having the formula:

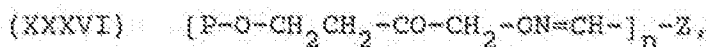


a compound having the formula:



and

a compound having the formula:

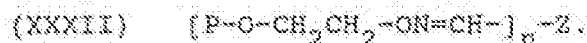


wherein Z is a polypeptide, n is 1 to x, x being the number of oxidation activatable groups on Z, and P is a water-soluble polymer.

5 25. A compound according to Claim 24, having the formula



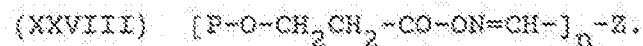
26. A compound according to Claim 24, having the formula



27. A compound according to Claim 24, having the formula

10 (XXXIII) $[P-O-CH_2CH_2-NH-CO-CH_2-ON=CH-]_n-Z$.

28. A compound according to Claim 24, having the formula



29. A compound according to any one of Claims 24-28, wherein n is 3 to 36.

15 30. A compound according to Claim 29, wherein n is 8-31.

31. A compound according to any one of Claims 7 to 17 or 24 to 30, wherein Z is selected from the group consisting of hormones, lymphokines, cytokines, growth factors, enzymes, vaccine antigens, and antibodies.

20 32. A compound according to Claim 31, wherein Z is a glycoprotein.

33. A compound according to Claim 32, wherein said glycoprotein is erythropoietin.

25 34. A compound according to any one of Claims 1 to 33, wherein said polymer P is selected from the group consisting of polyethylene glycol homopolymers, polypropylene glycol homopolymers, copolymers of ethylene glycol with propylene glycol, wherein said homopolymers

and copolymers are unsubstituted or substituted at one end with an alkyl group, polyoxyethylated polyols, polyvinyl alcohol, polysaccharides, polyvinyl ethyl ethers, α, β -poly[(2-hydroxyethyl)-DL-aspartamide], RO-PEG, where R may be alkyl, aryl, alkylaryl, aroyl, alkanoyl, benzoyl, arylalkylethers, cycloalkyl, cycloalkylaryl, and derivatives of said polymers.

5

35. A compound according to Claim 34, wherein said water-soluble polymer is polyethylene glycol or a derivative thereof.

10

36. A compound according to Claim 35, wherein said water-soluble polymer is monomethoxypoly(ethylene glycol).

37. A compound according to Claim 36, wherein the average molecular weight of the monomethoxypoly(ethylene glycol) is in the range of 2000-12000.

15

38. A compound according to Claim 37, wherein the average molecular weight of the monomethoxypoly(ethylene glycol) is 5000.

39. A water-soluble polymer modified polypeptide, said modified polypeptide produced by a method comprising the step of mixing a compound according to any one of Claims 1 to 6, 18 to 23 or any one of claims 34 to 38 as dependent on any one of claims 1 to 6 or 18 to 23, with a polypeptide for modification.

20

40. A modified polypeptide according to Claim 39, wherein said polypeptide for modification is selected from the group consisting of hormones, lymphokines, cytokines, growth factors, enzymes, vaccine antigens, and antibodies.

25

41. A modified polypeptide according to Claim 40,
wherein said polypeptide for modification is an antibody,
said method further comprising the step of combining said
antibody with a compound capable of specifically binding
5 to a binding site on said antibody, prior to said mixing
step.

42. A modified polypeptide according to Claim 40,
wherein said polypeptide for modification is an enzyme,
said method further comprising the step of combining said
10 polypeptide with a substrate for said enzyme, prior to
said mixing step.

43. A modified polypeptide according to Claim 40,
wherein said polypeptide for modification is a
glycoprotein.

15 44. A modified polypeptide according to Claim 43,
wherein said glycoprotein is erythropoietin.

45. A modified polypeptide according to any one of
Claims 39 to 44, said method further comprising the step
of adding an oxidizing agent prior to said mixing step.

20 46. A composition comprising a polypeptide according to
any one of Claims 39 to 45 and a pharmaceutically
acceptable carrier.

47. A method of activating polypeptides for conjugation
with compounds selected from the group consisting of
25 compounds II, III, IV, V, XIX, XXI, XXIII and XXIV said
method comprising the step of mixing a polypeptide for
activation with an oxidizing agent.

48. A method according to Claim 47, wherein said

oxidizing agent is sodium periodate.

49. A method according to Claim 48, wherein said periodate is present in a concentration of 10-40 micromoles per milligram of protein.

5 50. A method according to any one of Claims 47 to 49, wherein said mixing step takes place at temperature in the range of -10-50°C.

51. A method according to Claim 50, wherein said mixing step takes place at temperature in the range of 0-30°C.

10 52. A method according to any one of Claims 47 to 51, wherein said mixing step takes for a period of time between 1 minute and 3 days.

53. A method according to Claim 52, wherein said mixing step takes place for a period of time between 1 minute
15 and 60 minutes.

54. A method of making a water-soluble polymer modified polypeptide, said method comprising the step of mixing a water-soluble polymer reagent compound according to any one of Claims 1 to 6, 18 to 23 or any one of Claims 34 to
20 38 as dependent on any one of Claims 1 to 6 or 18 to 23, with a polypeptide for modification.

55. A method according to Claim 54, wherein said polypeptide for modification is selected from the group consisting of hormones, lymphokines, cytokines, growth
25 factors, enzymes, vaccine antigens, and antibodies.

56. A method according to Claim 55, wherein said polypeptide for modification is a glycoprotein.

57. A method according to Claim 56, wherein said glycoprotein is erythropoietin.

58. A method according to any one of Claims 54 to 57, said method further comprising the step of adding an oxidizing agent to the polypeptide for modification prior to said mixing step.

5 59. A method according to Claim 58, wherein said oxidizing agent is sodium periodate in a concentration in the range of 10-40 micromolar, P has a molecular weight in the range of 4000-12000, said water soluble polymer is the compound of formula

10 (III) $P-NH-CO-NHNH_2$.

60. A method according to Claim 58, wherein said oxidizing agent is sodium periodate in a concentration in the range of 10-40 micromolar, P has a molecular weight in the range of 4000-12000, said water-soluble polymer is
15 the compound of formula

(XXI) $P-O-CH_2CH_2-NH-CO-ONH_2$.

61. A kit for modifying polypeptides with water-soluble polymers, said kit comprising a water-soluble polymer according to any one of Claims 1 to 6 or 18 to 23 or any
20 one of claims 34 to 38 as dependent on any one of claims 1 to 6 or 18 to 23.

62. A kit according to Claim 61, said kit further comprising, an oxidizing agent.

63. A kit according to Claim 61 or 62, said kit further
25 comprising, a polypeptide for modification.

DATED this 8th Day of December, 1993

ORTEO PHARMACEUTICAL CORPORATION

Attorney: IAN ERNST
Fellow Institute of Patent Attorneys of Australia
of SHELSTON WATERS

ABSTRACT

The present invention provides methods and compounds for modifying polypeptides with PEG or other water-soluble organic polymers. Novel water-soluble polymer reagents are provided for.

The water-soluble polymer reagents of the subject invention include hydrazine, hydrazine carboxylate, semicarbazide, thiosemicarbazide, carbonic acid dihydrazide, carbazide, thiocarbazide, and arylhydrazide derivatives as well as oxylamine derivatives of water-soluble organic polymers, such as polyethylene glycol, polypropylene glycol, polyoxyethylated polyol, heparin, heparin fragments, dextran, polysaccharides, polyamino acids, and polyvinyl alcohol.

Also provided for, are polypeptides of interest derivatized by the subject water-soluble polymer reagents.

Kits for modifying polypeptides with the subject water-soluble polymer reagents are also provided.

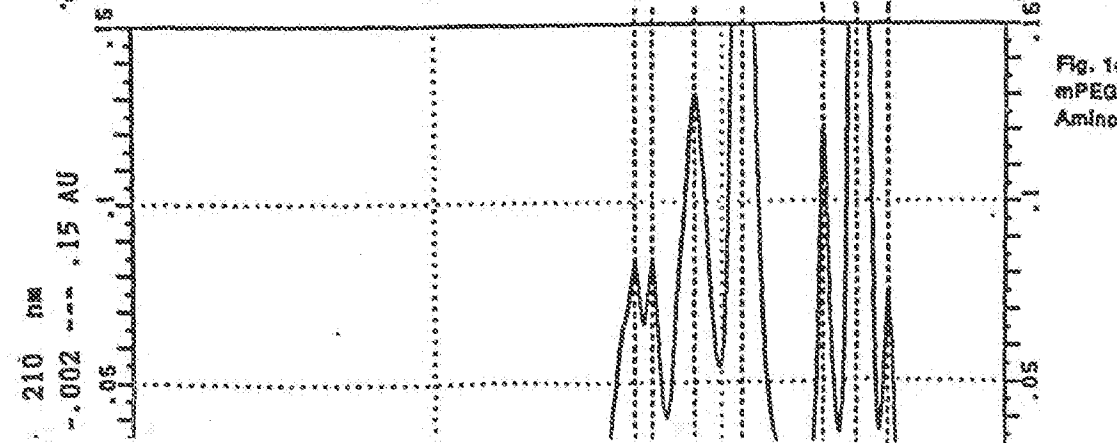
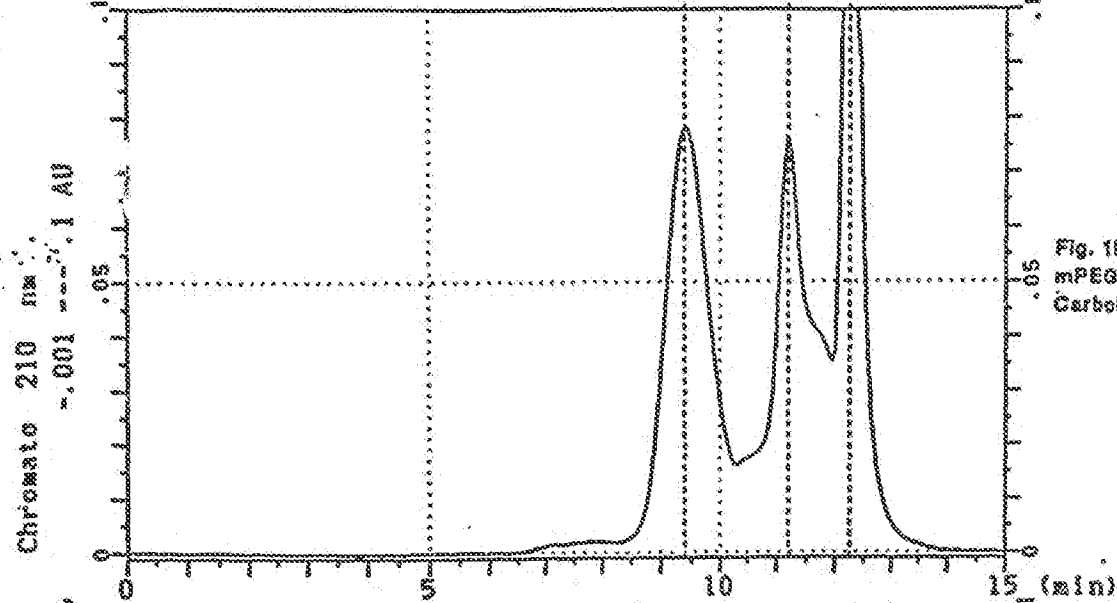
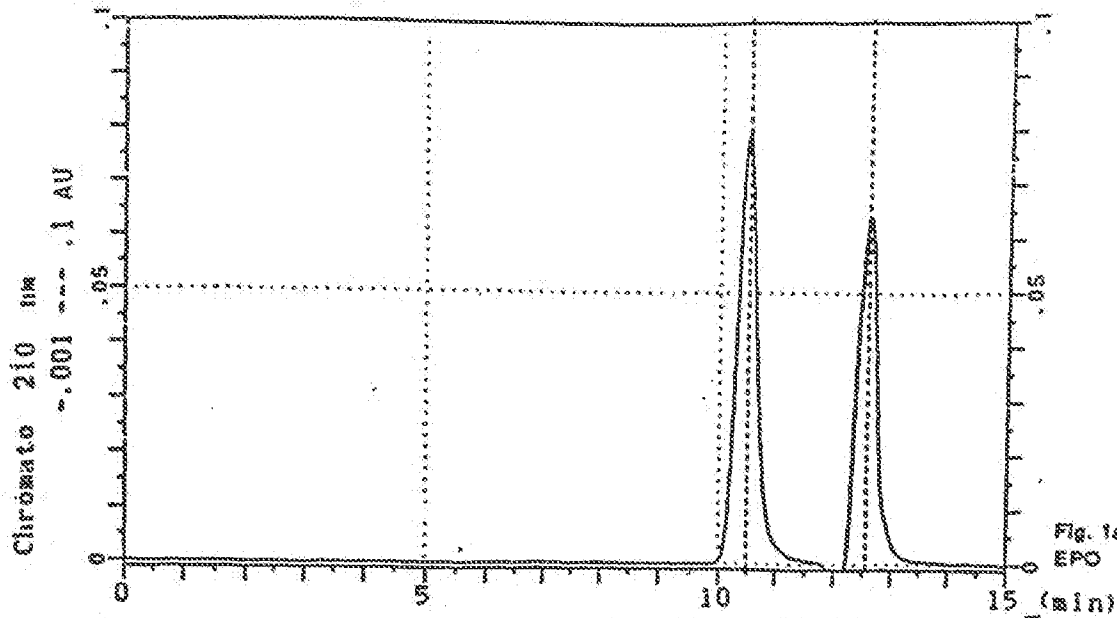
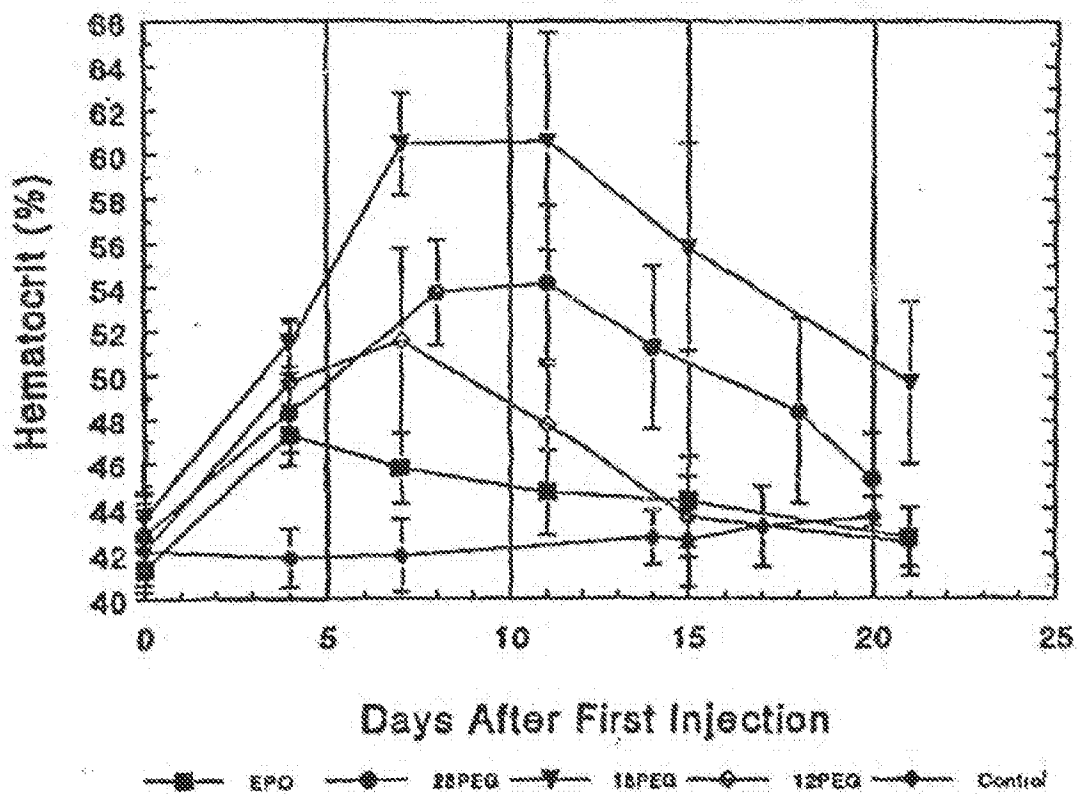


Figure 2: Hematocrit Levels in Mice
EPO and mPEG5000-EPOs,



EPO ELISA Assay

EPO and mPEG-EPOs

Fig. 3

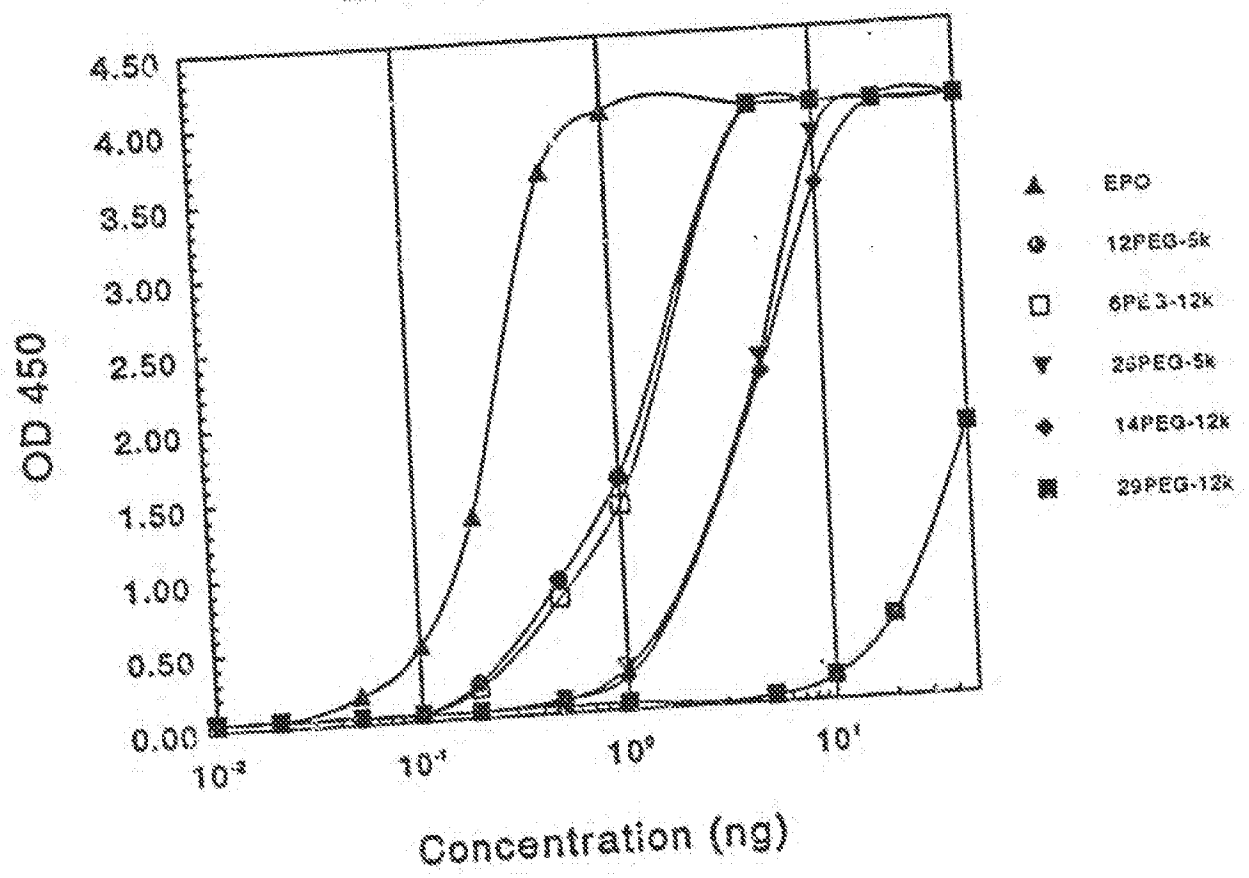


Figure 4: Hematocrit Levels in Mice
Hydrazide and Semicarbazide Comparison

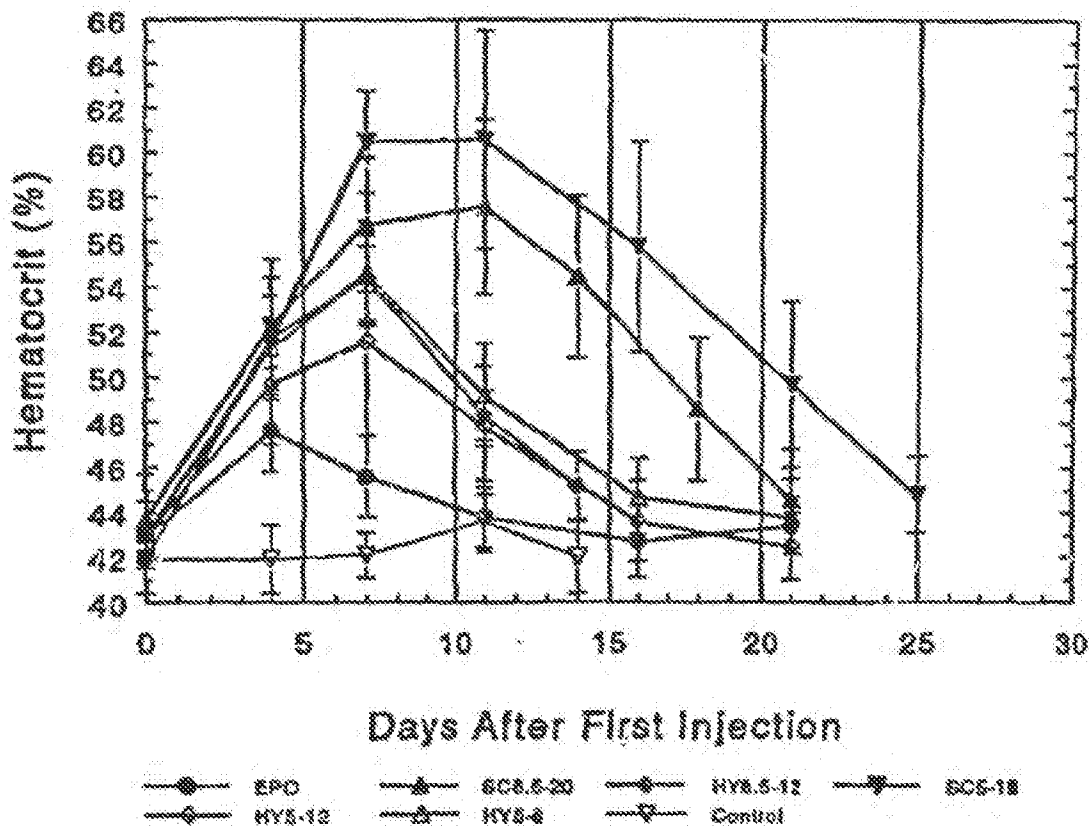


Figure 5: Hematocrit Levels in Mice
EPO and mPEG8500-EPOs,

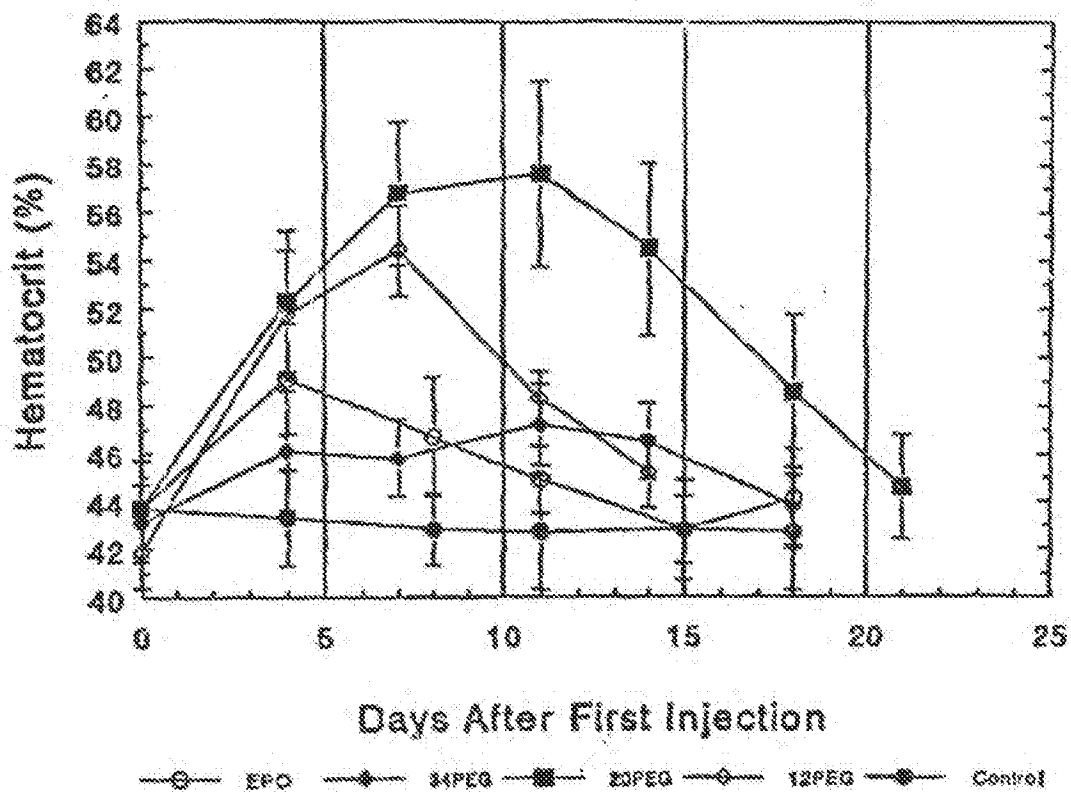
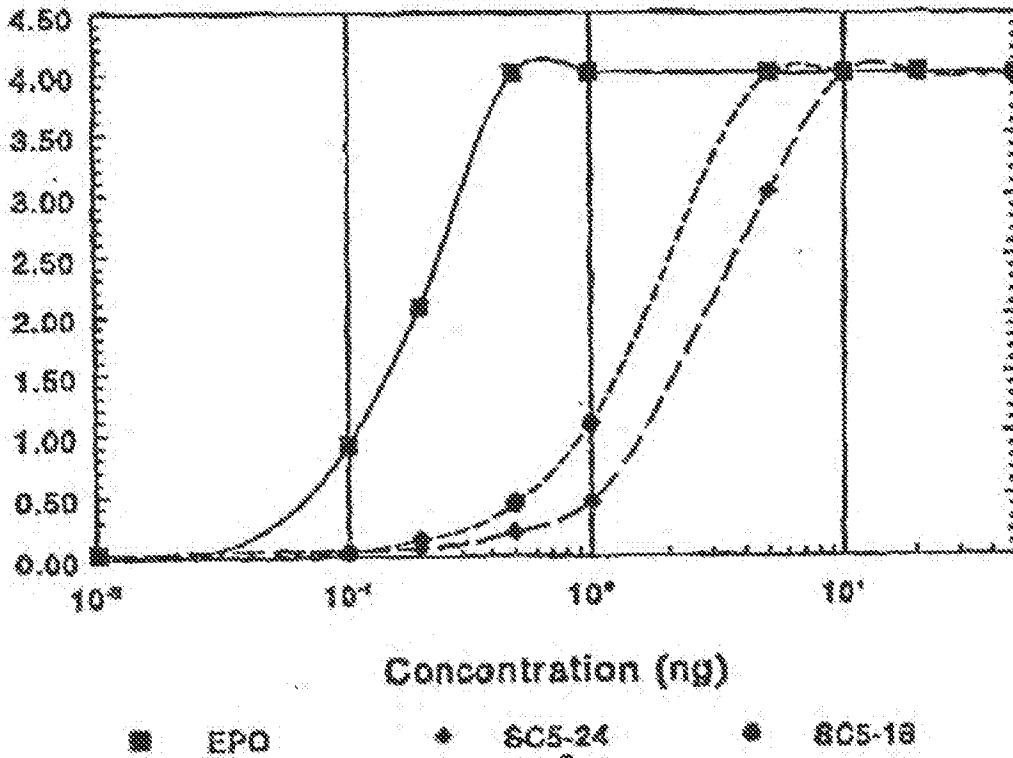


FIGURE 6

EPO ELISA Assay Carbohydrate Modification: Semicarbazide



Circulating Half-Life in Rat EPO Activity in Rat Plasma

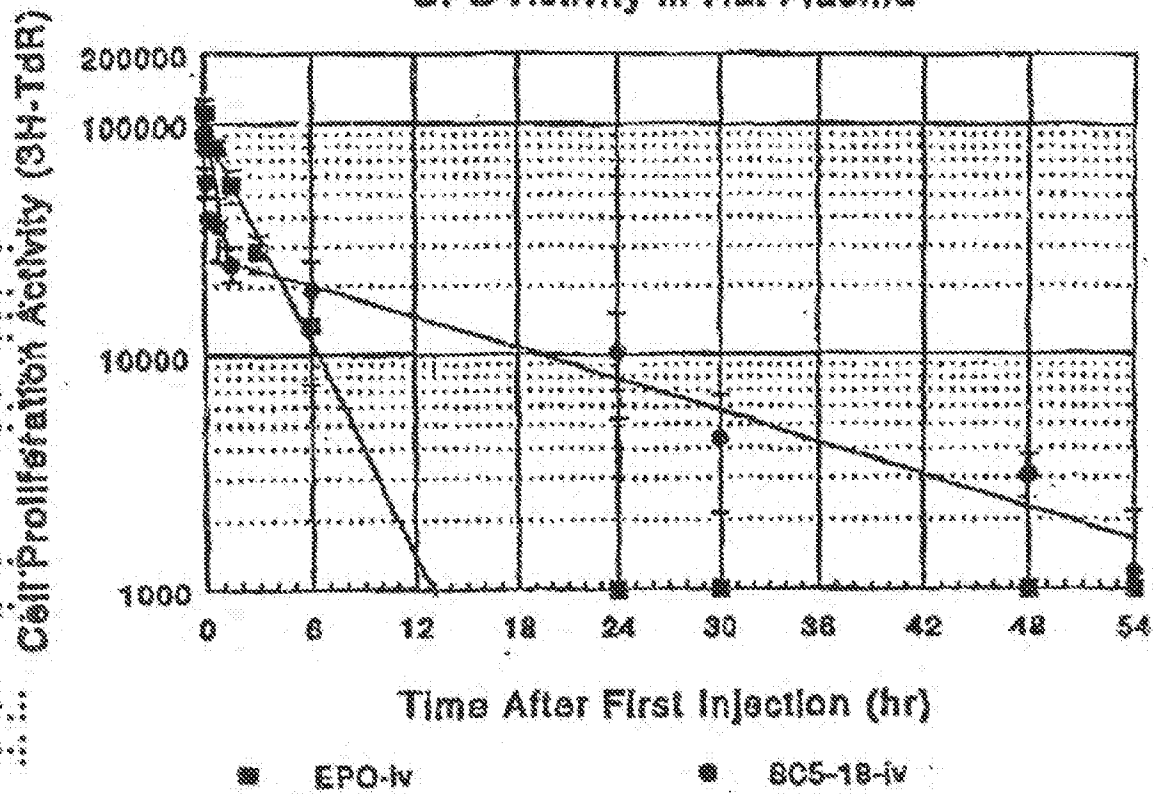


FIGURE 8

Hematocrit Levels in Normal Mice (CD1) 0.4ug / Dose Injected SC Days 0 & 1

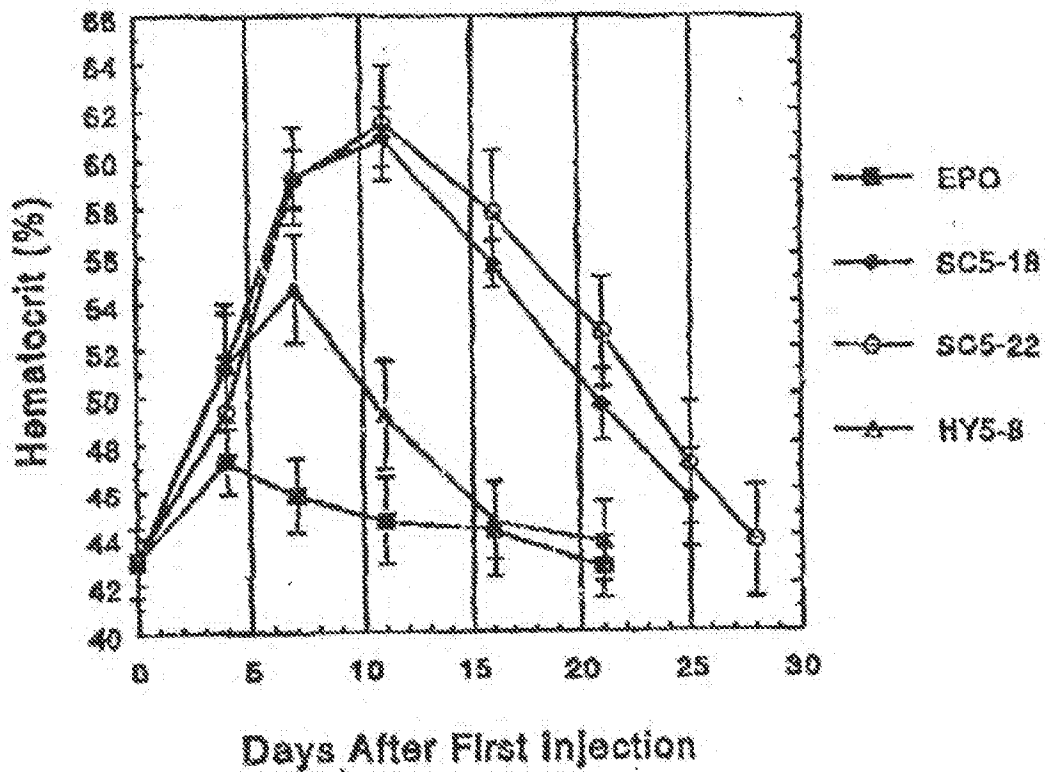
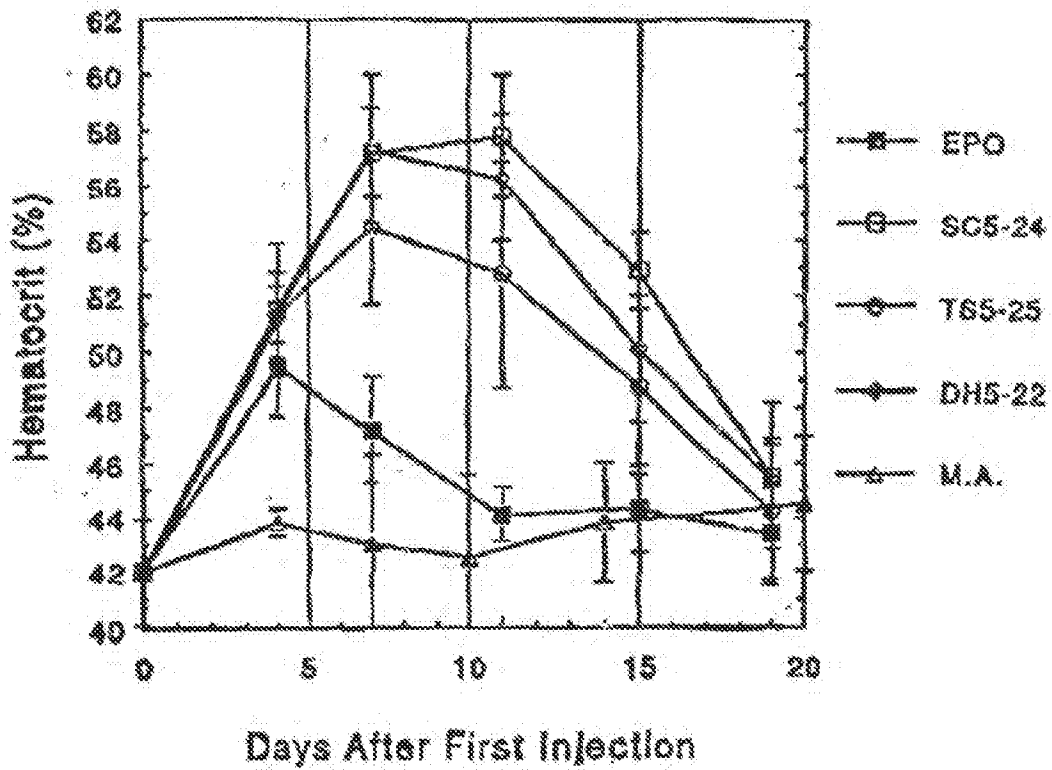
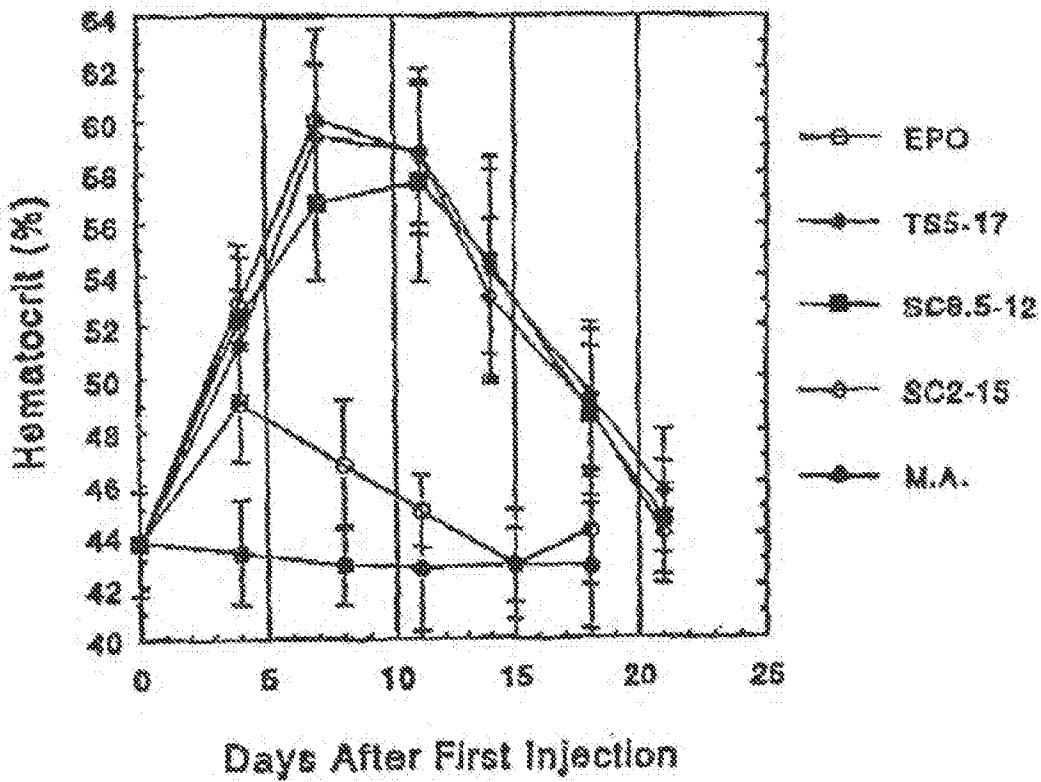


FIGURE 9

Hematocrit Levels in Normal Mice (CD1) 0.4ug / Dose Injected SC Days 0 & 1



Hematocrit Levels in Normal Mice (CD1) 0.4 ug/Dose Injected SC Days 0 & 1



Hematocrit Levels in Normal CD1 Mice
 0.4 ug / Dose Injected SC Days 0 & 1

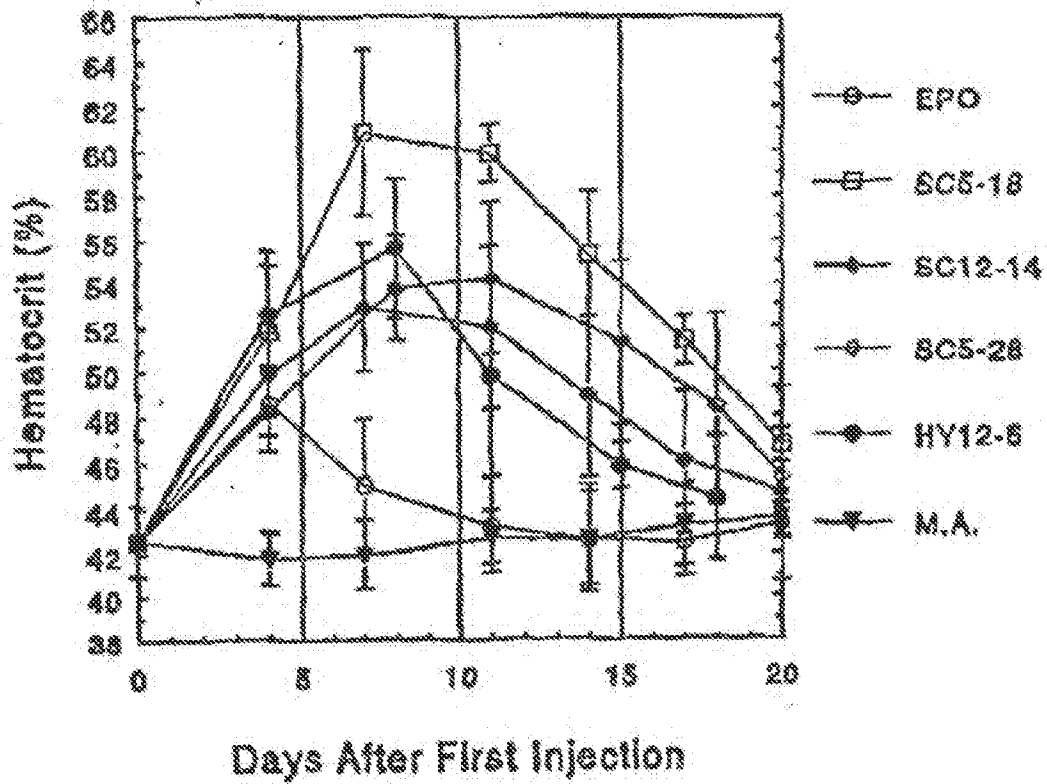
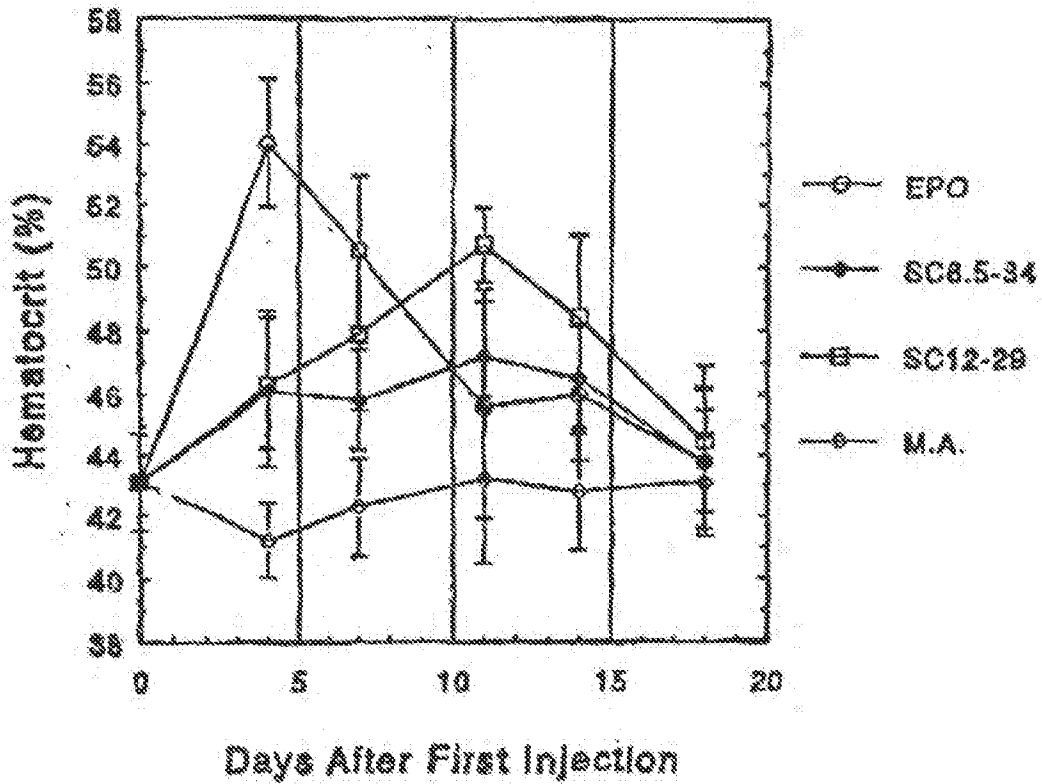
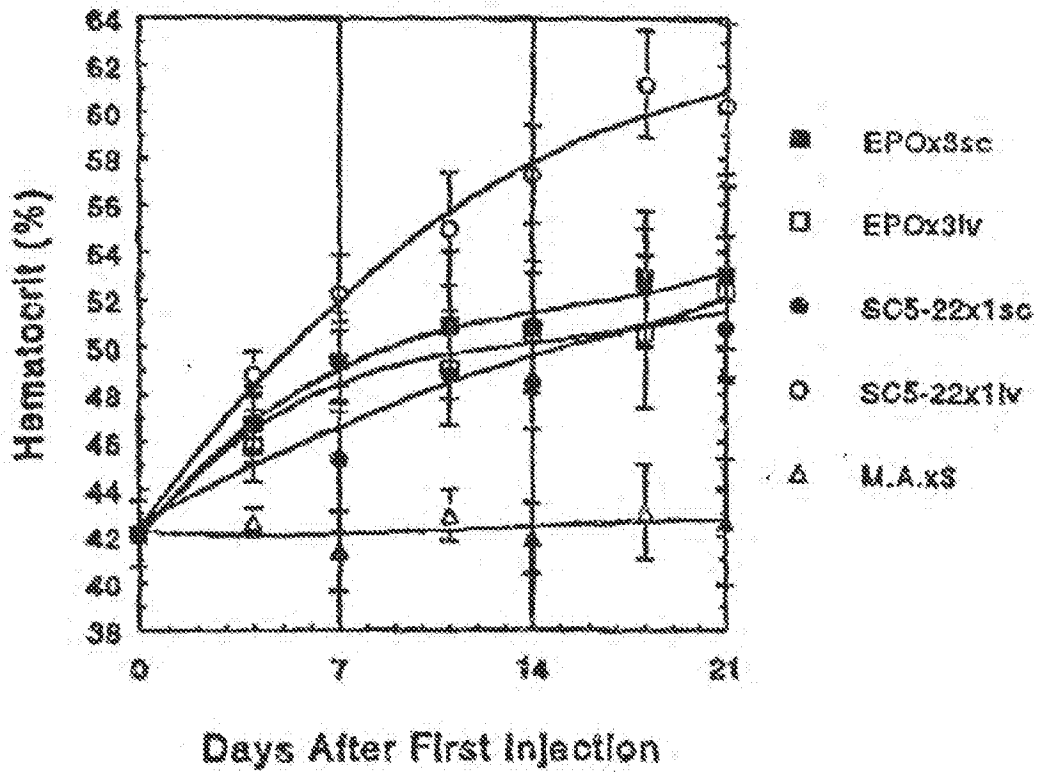


FIGURE 12

Hematocrit Levels in Normal CD1 Mice 0.4 ug / Dose Injected SC Days 0 & 1



Hematocrit Levels in Normal Mice (CD1) Multiple vs. Single Dose (0.1ug): SC/IV



Hematocrit Levels in Normal Mice (CD1) TNF α Induced Anemia

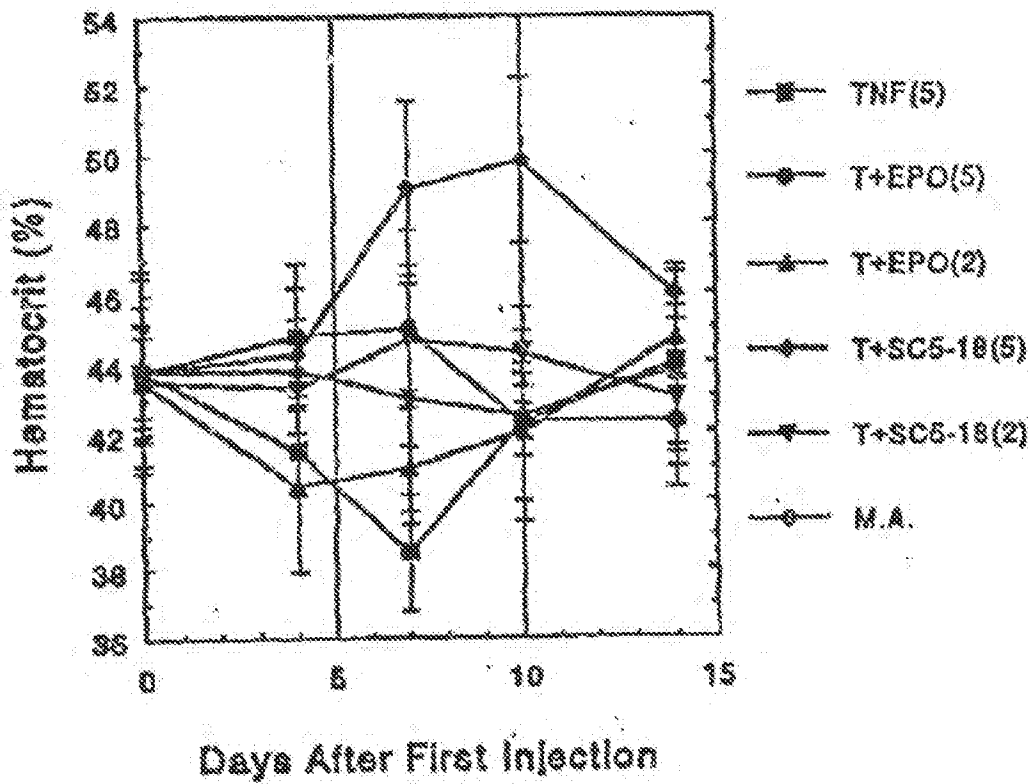


Figure 16

Hematocrit Levels in Normal Mice (CD1) 0.4ug / Dose Injected SC Days 0 & 1

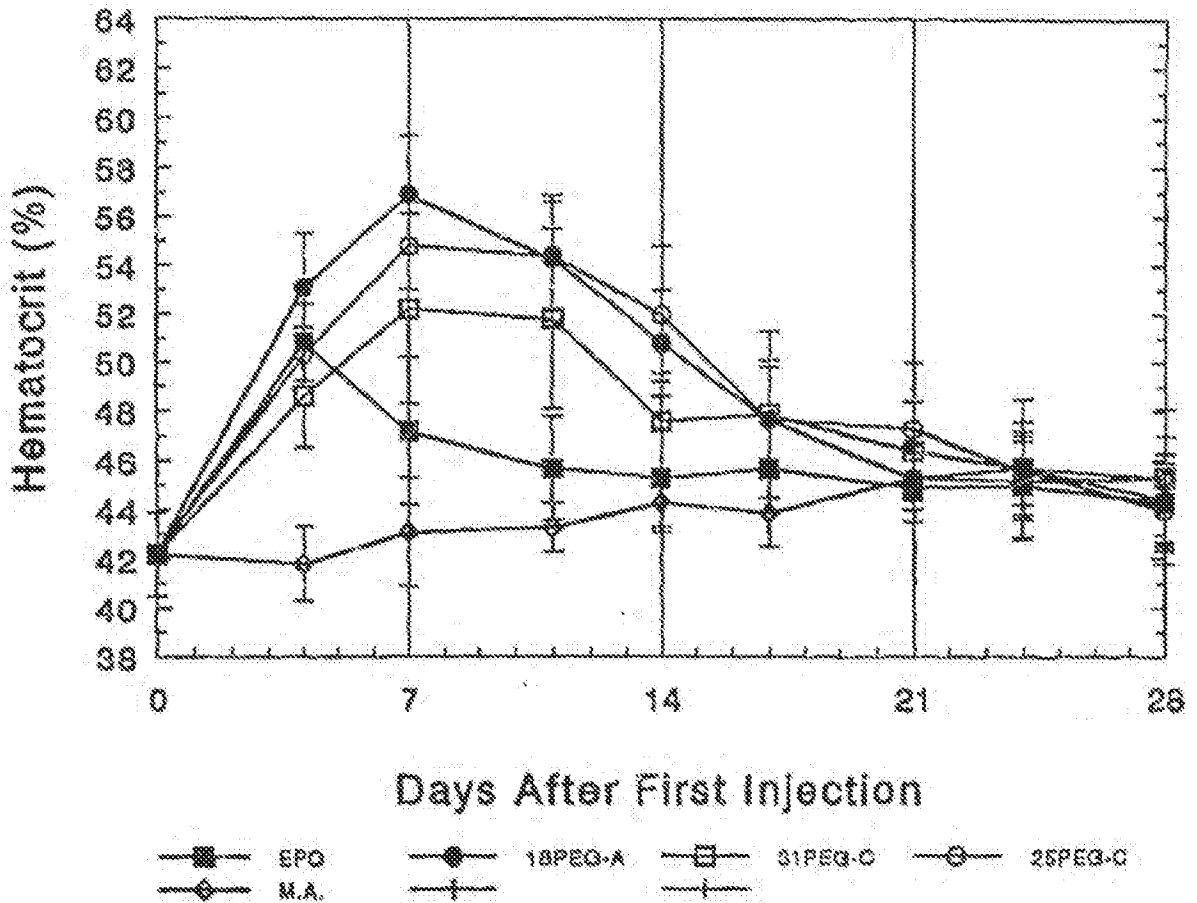


Figure 17

Hematocrit Levels in Normal Mice (CD1) 0.4ug / Dose Injected SC Days 0 & 1

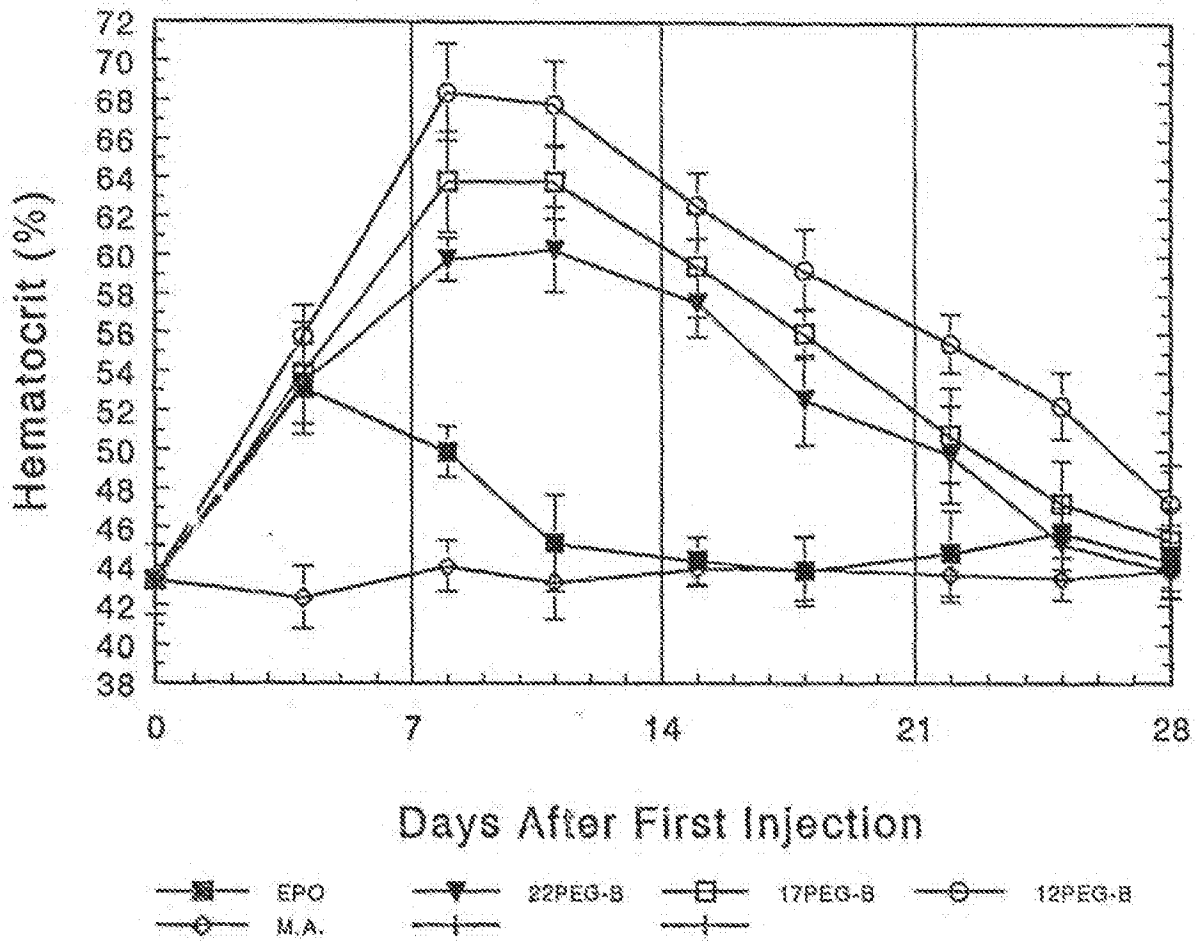


Figure 18

EPO ELISA Assay Oxime Linked mPEG-EPOs

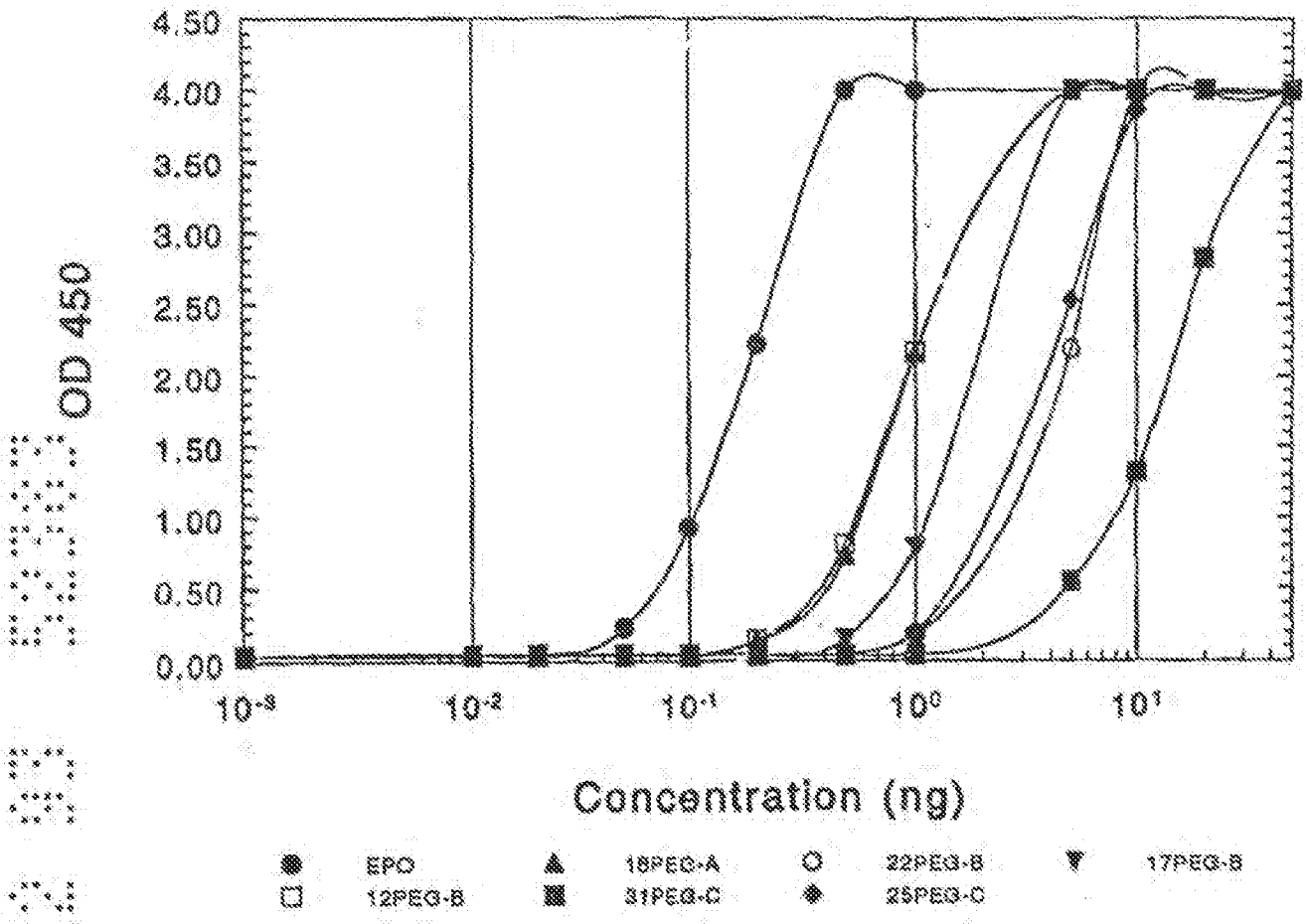
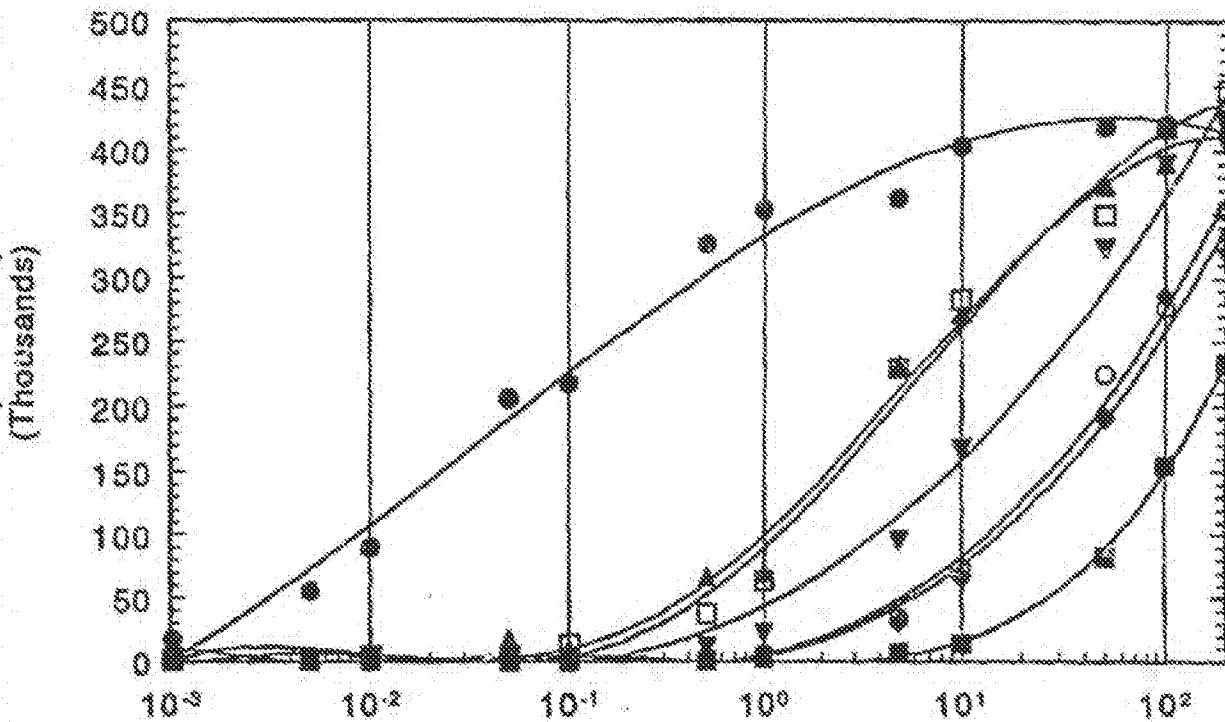


Figure 19

EPO Proliferation Assay Oxime Linked mPEG-EPOs

Proliferation Response (3H-TdR Incorporation)



Concentration (ng)

- EPO
- ▲ 18PEG-A
- 22PEG-B
- ▼ 17PEG-B
- 12PEG-B
- 31PEG-C
- ◆ 25PEG-C

Figure 20

Hematocrit Levels in Normal Mice (CD1) 0.4ug / Dose Injected SC Days 0 & 1

